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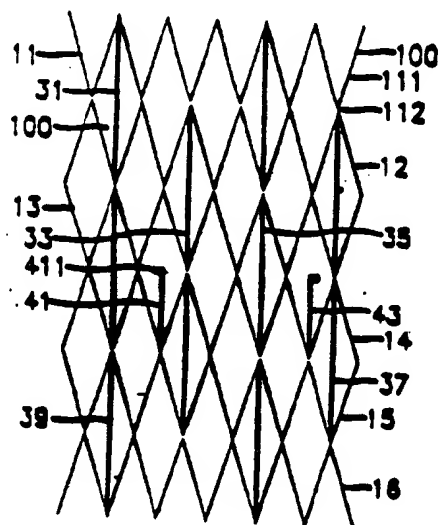
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(54) Title: SELF-EXPANDING ENDOVASCULAR STENT



(57) Abstract

An object of this invention is to provide a radially expandable stent (1) that holds a passageway enlarged by placing the stent (1) into a lumen. The stent (1) comprises a cylindrical frame formed by a plurality of unit structures (11, 12, 13, 31, 33, 35, 37, 39, 41, 43, 100, 111, 112, 12, 14, 15, 16...); said unit structures (11, 12, 13, 31, 33, 35, 37, 39, 41, 43, 100, 111, 112, 12, 14, 15, 16...) formed into a closed zig-zag configuration including an endless series of straight sections (111) and joined by bends (112), and arranged face to face into a shape of multilayer; connecting members (31, 33, 35, 37, 39, 41, 43) which connect said unit structures (11, 12, 13, 31, 33, 35, 37, 39, 41, 43, 100, 111, 112, 12, 14, 15, 16...); and a mesh (91), which is wrapped around an outside of said frame.

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SELF-EXPANDING ENDOVASCULAR STENT

Technical Field

5 This invention relates to improvements of
 endovascular stent

Background Art

10 It is desirable in various situations that means be
 provided for expanding a constricted vessel portion or
 for maintaining an open passageway through a vessel
 portion.

15 For example, these situations can be those by
 esophageal strictures that are caused by esophageal
 carcinoma or esophageal metastasis, or those by
 strictures that resulted from the cancer of biliary
 system, urinary duct system and bronchial system.

20 The balloon expansion has been well known method of
 enlarging and maintaining the passageway of esophagus in
 these cases, but this method should be operated
 repeatedly for its temporary effects, and also it has no
 effects on the patients of serious strictures.

25 As alternatives, various artificial-esophagi have
 been used in the cases of esophageal strictures, but
 since they have no constriction and relaxation, the rate
 of esophageal rupture is high (30 - 40%) in the process
 of inserting them into the strictural area that has
 been caused by the cancer, and the high mortality rate

due to mediastinitis results from rupture of esophagus. In addition, the patient has a great difficulty in swallowing due to narrow inner diameter (10 - 12 mm) of artificial esophagus, and obstruction of artificial-esophagus occurred frequently due to food intake.

As the means of overcoming the difficulty, a device to hold the passageway enlarged using a stent was presented by U.S. patent No. 4,214,587. However, the device of the invention has the temporary effect in enlarging the passageway, there is still the problem that the endovascular lumen gets narrows after a long time.

To improve this disadvantages, U.S. patent No. 4,580,568 was offered. The stent of said invention includes a wire formed into a closed zig-zag configuration including an endless series of straight sections joined by bends. The stent is resiliently compressible into a smaller first shape wherein the straight sections are arranged side-by-side and closely adjacent one another for insertion into a larger second shape wherein said straight sections press against the walls of the passageway to maintain it open.

However, in a case that the stent of the 4,580,568 invention is put into use for a long time, it still has the problem that the lumen is narrowed, because the proliferated cells cover the stent thickly. There is no effect in the case of the stricture by cancer, because the cancer cells can pass through the straight sections.

Disclosure of Invention

In view of the foregoing, it is the main object of this invention to provide a self-expanding endovascular stent, which expands a constricted vessel portion and maintains an open passageway through a vessel portion for a long time without moving in a lumen.

It is another object of the present invention to provide a self-expanding endovascular stent, which prevents the cancer cells to penetrate into a stent.

To accomplish said purposes, this invention provides, a stent, which comprises a cylindrical frame formed by a plurality of unit structures;

said unit structures formed into a closed zig-sag configuration including an endless series of straight sections and joined by bends, and arranged face to face into a shape of multistage; connecting members, which connects said unit structures one another and a mesh, which is wrapped around an outside of said frame.

And another stent according to the present invention comprises a cylindrical frame formed by a plurality of unit structures; said unit structures formed into a closed zig-sag configuration including an endless series of straight sections, joined by bends, and arranged face to face into a shape of multistage; connecting members, which connects said unit structures one another; anti-migration members, which have the same structure with said unit structure, and placed in the ends of the upper and lower portions of said frame; and

4

a mesh, which is wrapped around an outside of said frame.

Brief Description of Drawings

5

Fig. 1 is a development figure that illustrates a unit structure of a frame according to the present invention.

10 Fig. 2 is a perspective view of a first embodiment that illustrates a frame according to the present invention.

Fig. 3 is a development figure of said frame of the first embodiment of the invention.

15 Fig. 4 is a perspective view of the first embodiment according to the invention.

Fig. 5 is a view of showing a state that a stent of the first embodiment according to the present invention is placed and enlarged in a lumen.

20 Fig. 6 is a perspective view of a second embodiment according to the present invention that illustrates a frame.

25 Fig. 7 is a view of showing a state that a stent of the second embodiment of the present invention is placed and enlarged in a lumen.

Best Mode for Carrying out the Invention

For the purpose of promoting an understanding of

th principles of the invention, reference will now b
made to the embodiments illustrated in the drawings and
specific language will be used to describe the same. It
will nevertheless be understood that no limitation of
5 the scope of the invention is thereby intended, such
alterations and further modifications in the illustrated
device, and such applications of the principles of the
invention as illustrated therein being contemplated as
would normally occur of one skilled in the art to which
10 the invention relates.

Fig. 1 is a development figure that illustrates the
unit structure of a frame according to the present
invention; Fig. 2 is a perspective view of the first
embodiment that illustrates a frame according to the
15 present invention; Fig. 3 is a development figure of
said frame of Fig. 2.

A frame 10 of the first embodiment comprises a
reasonable number of unit structures 11 as shown in Fig.
1. The unit structure 11 is constructed of a wire, and
20 the wire is preferably made of stainless steel. The
unit structure 11 is formed into a closed sig-sag
configuration, thereby forming a series of straight
sections 111 and bends 112 as shown in Fig. 1. Both
ends of the wire are joined to each other by welding,
25 thereby forming an endless series of straight sections
joined by bends.

Frame 10 is constructed in a way that a plurality
of said unit structures 11 closely similar to each
other and bends corresponding to each of the unit

structures are arranged face to face into a shape of a multistage. At this time, every unit structure is connected in tandem by a reasonable number of connecting members 31, 33, 35, 37, These connecting members are joined appropriately between the bends of appointed unit structure such that they prevent each unit structure from being separated. The most preferable joint position of the connecting members is to be placed diagonally, regarding the connecting member 31 as a fiducial point as shown in Fig. 3. Namely, it is preferable to have a construction 31, 33, 35, and 37 as can be seen in Fig. 3. Also, the connecting members are not connected with every other bend and the frame 10 wherein an appropriate number of connecting members in one unit structure 11 i. e. ten connecting members in the first embodiment of Fig. 3 are connected to each other, has a plurality of diamond-like spaces 100. The number of the unit structure or the connecting members do not have a special meaning, and may be varied according to a state of a patient and/or convenience of production.

The space areas receive the straight section 111 when the straight section 111 is shrinked, making an elasticity variation with the bend 112.

A completed frame 10 has anti-migration members 41, 43 for preventing migration of a stent in a lumen that are equipped with barbs that jut outside. The anti-migration members 41, 43 are properly arranged and connected with the bends which the connecting members

31, 33, 35, 37 are not connected. Therefore, the stent 1 of the present invention does not move in a lumen.

All the unit structures, the connecting members, and the anti-migration members are made of the same material like stainless steel, preferably are gilt.

The frame 10 is wrapped in a mesh 91, and upper and lower hem of the mesh 91 are folded towards inside, and both hems become respectively adhesive whereby a wire of the unit structure 11 is not exposed. It is preferable that the mesh is made of nylon and the nylon mesh is coated with silicon rubber.

The method of 587 U.S. patent or a method using a catheter in which the stent of the invention being compressed may be used in order to place the stent of the present invention at a position, namely a diseased part in a lumen.

The stent is fixed on inner walls of the lumen with the barb 411 of the anti-migration member, and though the stent is used for a long time, it does not migrate, and in addition thereto, pain that a patient may feel can be relieved, compared with conventional stent, because an outer wall of the stent 1 is wrapped with mesh 91. Furthermore, the silicon rubber coating on the mesh 91 can prevent that cancer cell and the like penetrate into the inside of the stent.

As a result of experimenting the first embodiment of this invention on patients with gastric cancer or esophageal cancer, following effects are obtained.

A. A patient with gastric cancer and esophageal

spreading of gastric cancer came to Hospital of Cheon
Buk Medical College on July, 1989. The symptoms of th
patient were improved after six times medicine of
anti-cancer. But the patent revisited to the medical
college for an appearance of a dysphagia.

On January 8, 1990, an obstructed area of distal
portion of esophagus due to spreading of gastric cancer,
was enlarged with an esophageal balloon of 20 mm in
diameter and 8 cm in length, and then a stent of 20 mm
in external diameter and 8 cm in length was intubated
into a strictural area of esophagus through a sheath of
10 mm.

After operation, the patient was able to eat liquid
food and solid food, and in esophagogram performed in
seven days after operation, a balium passed well through
esophagus without any undercurrent or obstruction, and
specific complications such as the migration of the
stent or the rupture of the esophagus, did not occur.

After that, the patient left the hospital, and died
of a generalized metastasis of gastric cancer on April,
1990. However, there was no problem with the function of
the stent until just before the patient's death.

8. A patient with gastric cancer and esophageal
metastasis came to Hospital of Cheon Buk Medical College
owing to dysphagia on March 13, 1990.

After an obstructed portion of esophagus was
enlarged with an esophageal balloon of 20 mm in diameter
and 8 cm in length, a stent of 20 mm in external

diameter and 10 cm in length was intubated into the distal portion of esophagus through the sheath of 10 mm in external diameter.

As a result of the operation, the patient was not able to eat even water, but after the stent was intubated into his lumen, the patient was able to taking the liquid food and solid food, and a barium passed well through without any undercurrent or obstruction in esophagogram performed in seven days after the operation. When endoscope is performed on March 27, 1990, the endoscope of 10 mm in external diameter was passed through without any difficulty.

And in following up examination of esophagogram performed on August 10, 1990, any specific complications like the moving or the obstruction of the stent did not occur, and in addition, the patient was able to eat the liquid food and solid food.

C. A patient with the esophageal carcinoma was hospitalized in the Hospital of Cheon Buk Medical College, and took radiotherapy and medicated anti-cancer medicine five times, and then symptoms were improved, but the patient revisited to the hospital due to the dysphagia.

After the obstructed area of the esophagus was dilated with the esophageal balloon of 20 mm in diameter and 8 cm in length, the stent of 20 mm in external diameter and 12 cm in length was intubated into the obstructed area of the esophagus through the 10 mm sheath.

As a result of the treatment, the patient was able to eat liquid food and solid food, barium passed well through without any undercurrent or obstruction in esophagogram, and any complication like the migration of the stent or the rupture of the esophagus did not occur.

As medical appliance of the present invention is proved by the experiment on the above, since the expansile force of the stent that is intubated into an appointed area and is kept, is excellent, the outstanding effects that keep the stricture of passageway enlarged can be obtained.

The construction of the first embodiment as mentioned above is expected of preventing the migration of the stent, but since the distal ends of the anti-migration members 41, 43 have barbs that jut out externally, the possibility cannot be excluded that the barbs cause a patient to feel pain and lumen perforation, and when the stent is artificially transferred to set the position of the stent to rights, the lumen wall can be hurt.

Fig. 6 is a perspective view of a second embodiment of the present invention, and shows that a plurality of unit structures 21, 22, 23, 24 are linker, being expanding. The unit structures 21, 22, 23, 24 are identical with the unit structures of the first embodiment.

According to the second embodiment of this invention, the stent comprises the frame 20 formed by a plurality of connected unit structures 21, 22, 23, 24,

and anti-migration members 42 and 44 placed in lower and upper portions of the frame 20. The frame 20 without anti-migration members 42 and 44 is almost same with the frame of the first embodiment of this invention except for the numbers of the unit structures and the length of connecting members 36, 38. The numbers of unit structures have no meaning in this invention. And the length of connecting members 36, 38 in the ends of the upper and lower portions of the frame 20 is a half of the length of the connecting members 31, 39 of the first embodiment of this invention.

The both side sections of the unit structures 21, 24 are connected with the anti-migration members 42 and 44 through the second connecting members 32, 34.

Anti-migration members 42 and 44 placed in the ends of the upper and lower portions of the frame 20 have the same structures with the unit structures 21, 22, 23, 24. But the anti-migration members 42 and 44 are formed to be larger than the frame 20 in diameter. To connect the frame 20, and the anti-migration members 42 and 44, that are different from each other in diameter, is available through the second connecting members 32, 34. The second connecting member 34 connects a bend 381 of the end 381 of the connecting member 38 with a bend 442 of the anti-migration member 44. This connection is performed by welding.

Since the diameter of the anti-migration members 42 and 44 depends on the length of horizontal parts 321, 341 of the second connecting members 32, 34, it is

important to determine the length thereof. Expansile parts 322, 342, expanded from the horizontal parts 321, 341 and bended vertically.

5 It is omitted in Fig. 6 that a mesh is wrapped around the outside of the frame 20.

The stent 2 formed as mentioned above keeps expanded unless force is given thereto. In order to place the stent within an esophagus, a given introducing assembly such as an introducing tube, an esophageal balloon and a pusher catheter are pushed into a stricture in a compressed state.

10 As the introducing tube is removed over the pusher catheter, the compressed stent is automatically passed through and expanded.

15 Fig. 7 is an explanatory view of the state that the stent 2 placed within a lumen is radially expanded to stick to the lumen.

20 A stent that can be used for patients with esophageal carcinoma is described above, and if a stent was formed to be cylindrical between 8 mm to 10 mm in diameter, other applications of the stent are in the biliary, bronchial and urinary systems.

CLAIMS

1. A stent comprising:
a cylindrical frame 10 formed by a plurality of
5 unit structures 11, 12, 13, 14, 15, 16;
said unit structures 11, 12, 13, 14, 15, 16 formed
into a closed sig-sag configuration including an
endless series of straight sections 111 and joined by
bends 112, and arranged face to face into a shape of
10 multistage;
connecting members 31, 33, 35, 37, which connects
said unit structures 11, 12, 13, 14, 15, 16;
and a mesh 91, which is wrapped around an outside
of said frame 10.
- 15 2. A stent as claimed in claim 1 wherein
anti-migration members 41, 43 with barbs are installed
in a central area 100 of said frame 10.
3. A stent as claimed in claims 1 and 2 wherein
said frame 10, connecting members 31, 33, 35, 37, and
20 anti-migration members 41, 43 are gilt.
4. A stent as claimed in claims 1 and 2, wherein
said mesh 91 coated with silicon rubber.
5. A stent comprising:
a cylindrical frame 20 formed by a plurality of
25 unit structures 21, 22, 23, 24;
said unit structures 21, 22, 23, 24 formed into a
closed sig-sag configuration including an endless series
of straight sections 211, joined by bends 212, and
arranged face to face into a shape of multistage;

connecting members, which connects said unit structures 21, 22, 23, 24;

anti-migration members 42 and 44, which have same structures with said unit structure, and placed in the ends of the upper and lower portions of said frame; and a mesh, which wrapped around an outside of said frame 20.

6. A stent as claimed in claim in 5, wherein said anti-migration members 42 and 44 have larger diameters than the diameter of said frame 20, and connected to said frame 20 by the second connecting members 32, 34.

7. A stent as claimed in claim 6, wherein the second connecting members 32, 34 consisting of vertical parts 321, 341 and expansile parts 322, 342.

8. A stent as claimed in claims 5, 6 and 7, wherein a mesh wrapped whole around an outside of said frame 20 and anti-migration members 42 and 44.

9. A stent as claimed in claims 5 and 6, wherein said frame 20 and anti-migration members 42, 44 are gilt.

10. A stent as claimed in claims 5 and 8, wherein said mesh coated with silicon rubber.

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FIG. 1

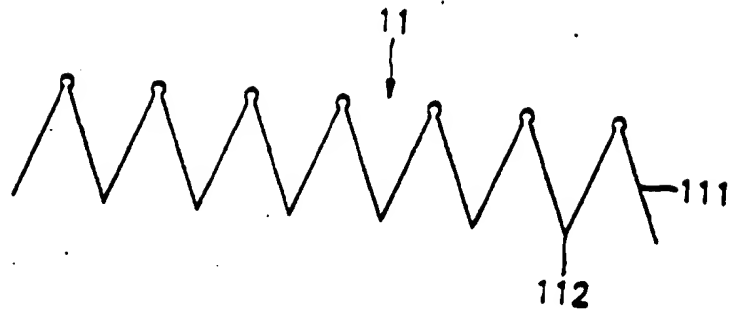
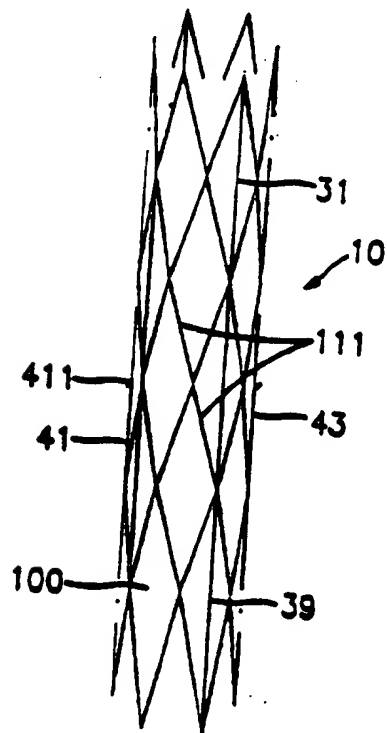


FIG. 2



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FIG. 3

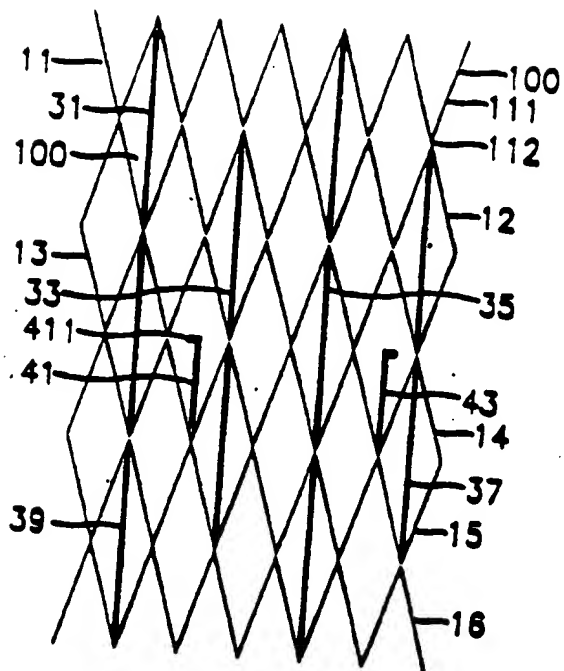
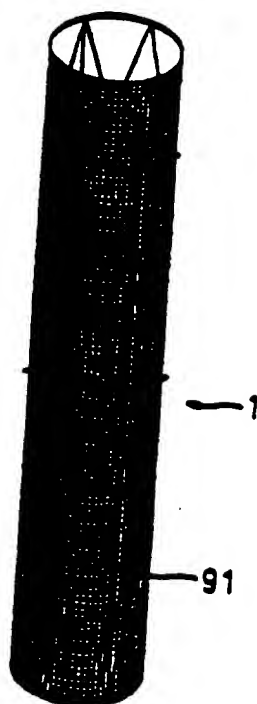


FIG. 4



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FIG.5

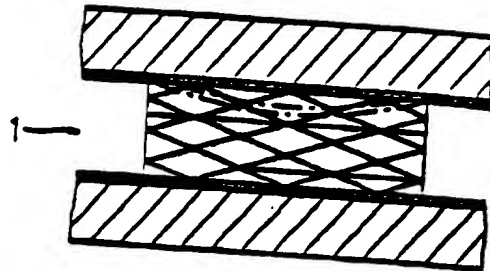


FIG.6

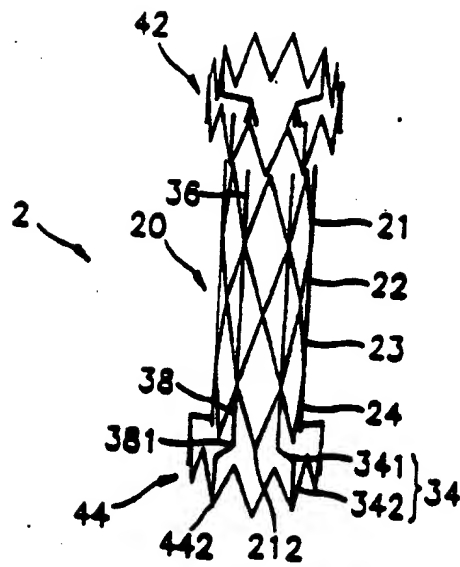
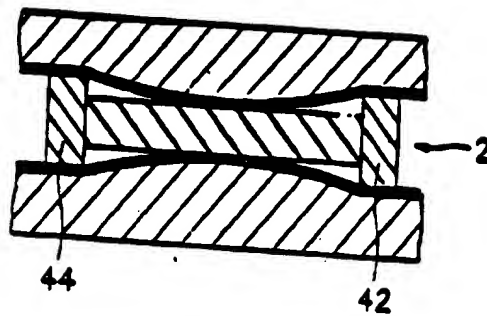


FIG.7



INTERNATIONAL SEARCH REPORT

International Application No. PCT/KR 91/00023

I. CLASSIFICATION OF SUBJECT MATTER (If several classification symbols apply, indicate all)

According to International Patent Classification (IPC) or to both National Classification and IPC

Int. Cl.⁵: A 61 M 29/00

II. FIELDS SEARCHED

Minimum Documentation Searched?

Classification System

Classification Symbols

Int. Cl.⁵: A 61 M 29/00; A 61 F 2/04; A 61 M 25/00

Documentation Searched other than Minimum Documentation
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III. DOCUMENTS CONSIDERED TO BE RELEVANT¹

Category² | Character of Document, ³ with indication, where appropriate, of the relevant passages ⁴ | Relevant to Claim No. ⁵

- | | | |
|---|---|-------|
| A | US, A, 4 733 665 (PALMAZ) 29 March 1988
(29.03.88), see especially figures 1A, 1B; column
5 line 24 to column 6, line 66. | (1,5) |
| A | DE, B, 1 766 921 (NAT. RES. DEV.) 15 January 1970
(15.01.70), see especially figures 1-4, 9;
claims 1, 2, 9, 11, 13, 14. | (1,5) |
| A | US, A, 4 580 568 (GIANTURCO) 08 April 1986
(08.04.86), see especially figures 1, 6-8; column 2,
lines 37 to 68; column 3, lines 17 to 25. | (1,5) |

¹ Special categories of cited documents:

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⁹ Y⁹ document of particular relevance: the claimed invention
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in the art

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IV. CERTIFICATION

Date of the Actual Completion of the International Search

17 January 1992 (17.01.92)

Date of Mailing of this International Search Report

24 January 1992 (24.01.92)

Signature of Authorized Officer

PCT/IR 91/00023

In diesem Anhang sind die Mitglieder
der Patentfamilien der in diesem
Anhang internationalen Recherchenbericht
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This Annex lists the patent family
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cited in the above-mentioned inter-
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